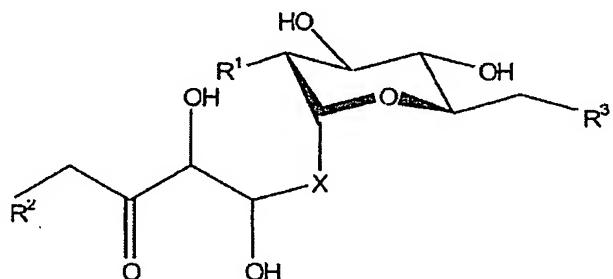


Amendments to the Claims:

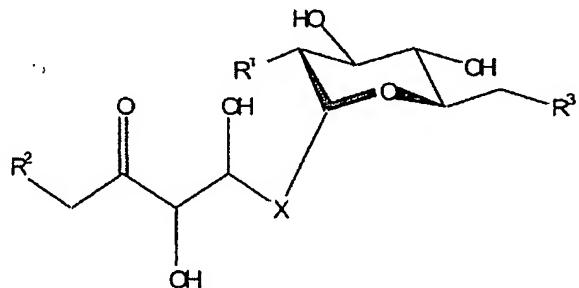
This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (currently amended) Compounds-A compound having affinity to and/or selectivity for P-selectin, represented by the following and having structure of formula Ia:



and their or a stereo-isomers thereof, represented by the following formula Ib:



wherein:

X is an optional group, which represents -O-, -OCH₂-, -S-, -SCH₂-, -NH- or -NHCH₂;

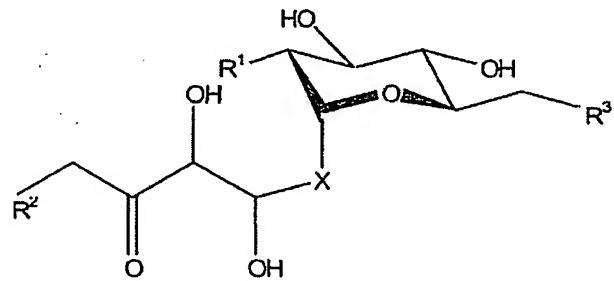
R¹ represents QR⁴, herein Q represents- O-, -NH-, -NH(C=O)-, -O(C=O), -NH(C=O)-O- or -NH(C=O)-NH; and wherein R⁴ represents any substituent comprising at least one carbon atom;

R² is a moiety bearing at least one negative charge and

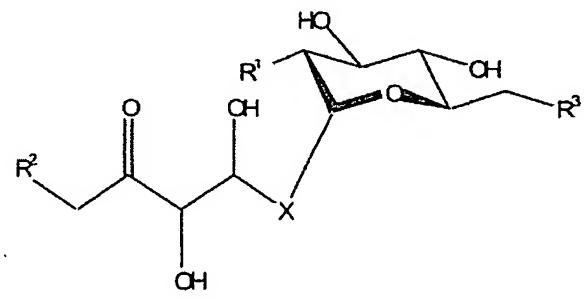
R³ can be is any group.

2. (currently amended) The compounds according to claim 1, wherein X is not present or represents -O-.
3. (currently amended) The compounds according to any one of the preceding claims claim 1, wherein Q represents -NH(C=O)-.

4. (currently amended) The compounds according to ~~any one of the preceding claims~~claim 1, wherein R² is or comprises a phosphate group.
5. (currently amended) The compounds according to ~~any one of the preceding claims~~claim 1, wherein R³ represents OH or YR⁵, wherein Y is -O-, -CH₂- or -NH- and R⁵ comprises at least one carbon atom.
6. (currently amended) The compounds according to ~~any one of the preceding claims~~claim 1, wherein R⁴ comprises an alkyl moiety, an aromatic moiety or a group comprising an electron withdrawing moiety.
7. (currently amended) The compounds according to claim 6, wherein R⁴ is a phenyl or a naphthalene group.
8. (currently amended) Compounds-A compound having affinity to and/or selectivity for P-selectin, represented by the formula Ia,



and ~~their~~ or a stereo-isomers thereof, represented by the formula Ib,



-wherein:

X is an optional group, which represents -O-, -OCH₂-, -S-, -SCH₂-, -NH- or -NHCH₂;

R¹ represents QR⁴, herein Q represents -O-, -NH-, -NH(C=O)-, -O-(C=O), -NH-(C=O)-O- or -NH-C=O)-NH-; and wherein R⁴ represents any substituent comprising at least one carbon atom;

R² is a moiety bearing at least one negative charge and

R^3 can be is any group,

wherein R³ comprises an anchor moiety capable of anchoring the compound to a colloidal or microparticulate drug carrier.

9. (currently amended) The compounds according to claim 8, wherein the anchor moiety is a peptide or peptidomimetic moiety having affinity to P-selectin.

10 - 13 (canceled)

14. (currently amended) Pharmaceutical-A composition, comprising in a pharmaceutically acceptable carrier a compound according to claim 1 having affinity to and/or selectivity for P-selectin, represented by the formula Ia, and its stereoisomer, represented by the formula Ib, wherein:

X is an optional group, which represents O, OCH₂, S, SCH₂, NH or NHCH₂;

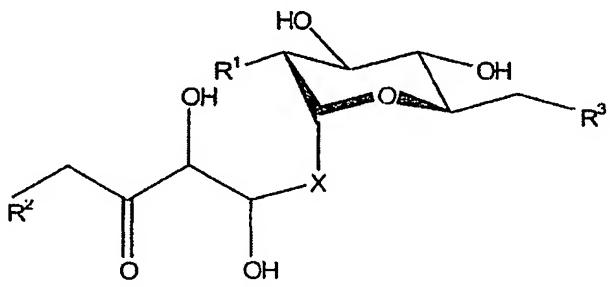
R¹ represents QR⁴, herein Q represents O, NH, NH(C=O), O(C=O), NH(C=O)O or NH(C=O)NH; and wherein R⁴ represents any substituent comprising at least one carbon atom;

R² is a moiety bearing at least one negative charge and

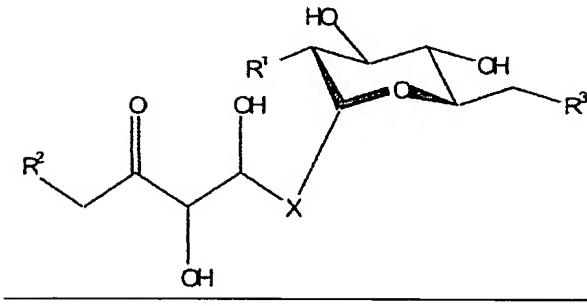
R³ can be any group,

or a derivative, salt, conjugate, solvate, or multimer thereof

15. (currently amended) A method for determining whether a compound is capable of binding to P-selectin or a functional equivalent of P-selectin, comprising contacting and incubating the compound to be tested and a predetermined amount of a compound having affinity to and/or selectivity for P-selectin, represented by the formula Ia,



and its or a stereo-isomer thereof, represented by the formula I b-,



wherein:

X is an optional group, which represents -O-, -OCH₂-, -S-, -SCH₂-, -NH- or -NHCH₂-;

R¹ represents QR⁴, herein Q represents -O-, -NH-, -NH(C=O)-, -O-(C=O), -NH-(C=O)-O- or -NH-(C=O)-NH-; and wherein R⁴ represents any substituent comprising at least one carbon atom;

R² is a moiety bearing at least one negative charge and

R' can be any group,

with a predetermined amount of P-selectin or said functional equivalent of P-selectin and

subsequently determining the amount of the same compound.

16. (new) A method of treating or inhibiting a disease or condition involving activation and/or overexpression of P-selectin in a mammal inflicted with such a disease, the method comprising administering to the mammal an effective P-selectin inhibiting amount of a composition according to claim 14.